SCORE Search Results Details for Application 10568337 and Search Result 20071129_084935_20071129_084935_us-10-568-337-2.p2n.rni.

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<u>Overview</u>

SCORE <u>FAQ</u>

Comments / Suggestions

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OM protein - nucleic search, using frame plus p2n model

November 29, 2007, 08:49:46; Search time 836 Seconds Run on:

(without alignments)

120.985 Million cell updates/sec

Title: US-10-568-337-2

Perfect score: 124

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Searched: 5155175 segs, 1873024446 residues

Total number of hits satisfying chosen parameters: 10310228

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 1000 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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	3	55	44.4	36	2	US-07-731-157A-11	Sequence 11, Appl	
	4	55	44.4	36	2	US-08-541-780-11	Sequence 11, Appl	
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С	6	54	43.5	539	5	US-10-703-032-91504	Sequence 91504, A	
С	7	54	43.5	588	5	US-10-703-032-94706	Sequence 94706, A	
С	8	53	42.7	419	5	US-10-703-032-49153	Sequence 49153, A	
С	9	52.5	42.3	378	5	US-10-703-032-89469	Sequence 89469, A	
С	10	52.5	42.3	396	5	US-10-703-032-98346	Sequence 98346, A	
С	11	52.5	42.3	418	5	US-10-703-032-98096	Sequence 98096, A	
С	12	52.5	42.3	422	5	US-10-703-032 - 96699	Sequence 96699, A	
C	13	52.5	42.3	425	5	US-10-703-032-97933	Sequence 97933, A	
С	14	52.5	42.3	428	5	US-10-703-032-80385	Sequence 80385, A	
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; Patent No. 5457032
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    APPLICANT: Misset, Onno
    APPLICANT: Van der Laan, Jan M.
    APPLICANT: Lenting, Herman B.M.
    TITLE OF INVENTION: Mutated beta-lactam acylase genes
    NUMBER OF SEQUENCES: 50
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: COOLEY GODWARD CASTRO HUDDLESON & TATUM
      STREET: FIVE PALO ALTO SQUARE, 4TH FLOOR
      CITY: PALO ALTO
      STATE: CALIFORNIA
      COUNTRY: USA
      ZIP: 94306
    COMPUTER READABLE FORM:
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      APPLICATION NUMBER: US/07/731,157A
      FILING DATE: 19910509
      CLASSIFICATION: 435
    PRIOR APPLICATION DATA:
      APPLICATION NUMBER: EP 90200962
      FILING DATE: 18-APR-1990
    ATTORNEY/AGENT INFORMATION:
    NAME: RAE-VENTER PH.D., BARBARA
      REGISTRATION NUMBER: 32,750
      REFERENCE/DOCKET NUMBER: GBRO-027/00US
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: 415-494-7622
       TELEFAX: 415-857-0663
      TELEX: 380816 COOLEY PA
   INFORMATION FOR SEQ ID NO: 1:
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; Sequence 1, Application US/08541780
; Patent No. 5935831
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    APPLICANT: Quax, Wilhelmus J.
    APPLICANT: Misset, Onno
;
    APPLICANT: Van der Laan, Jan M.
    APPLICANT: Lenting, Herman B.M.
    TITLE OF INVENTION: Mutated beta-lactam acylase genes
    NUMBER OF SEQUENCES: 50
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: COOLEY GODWARD CASTRO HUDDLESON & TATUM
      STREET: FIVE PALO ALTO SQUARE, 4TH FLOOR
      CITY: PALO ALTO
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      APPLICATION NUMBER: US/07/731,157
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    ATTORNEY/AGENT INFORMATION:
      NAME: RAE-VENTER PH.D., BARBARA
      REGISTRATION NUMBER: 32,750
     REFERENCE/DOCKET NUMBER: GBRO-027/00US
    TELECOMMUNICATION INFORMATION:
      TELEPHONE: 415-494-7622
      TELEFAX: 415-857-0663
      TELEX: 380816 COOLEY PA
  INFORMATION FOR SEQ ID NO: 1:
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      TYPE: nucleic acid
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    HYPOTHETICAL: NO
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	SCORE Search Results Details for Application 10568337 and Search Result 20071129_0	Page 4 of 4
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	•	
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SCORE Search Results Details for Application 10568337 and Search Result 20071128_153802_us-10-568-337-5.rng.

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SCORE System **Overview**

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GenCore version 6.2.1 Copyright (c) 1993 - 2007 Biocceleration Ltd.

OM nucleic - nucleic search, using sw model

Run on:

November 29, 2007, 00:22:26; Search time 646 Seconds

(without alignments)

1147.147 Million cell updates/sec

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US-10-568-337-5

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Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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	8	94.8	94.8	101	5	AAH27740	Aah27740 Gl-7ACA r
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AC
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DT
    05-MAY-2005 (first entry)
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XX
KW
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KW
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XX
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XX
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PN
XX
PD
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XX
PF
    12-AUG-2004; 2004WO-EP009055.
XX
    13-AUG-2003; 2003US-0494915P.
PR
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PΑ
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XX
PΙ
    Stempfer G, Alliger P, Palma N;
XX
    WPI; 2005-182386/19.
DR
XX
    Preparing recombinant polypeptides of interest, for producing large
PT
    variety of polypeptides of interest, by fermenting prokaryotic host cell
PT
PT
    comprising a periplasm transformed with a recombinant expression system.
XX
PS
    Example 1; SEQ ID NO 8; 29pp; English.
XX
CC
    This invention relates to a novel method of preparing a recombinant
CC
    polypeptide of interest which comprises fermenting a prokaryotic host
    cell comprising a periplasm transformed with a recombinant expression
CC
CC
    system capable of bringing secretion of a polypeptide of interest into
CC
    the periplasm of the host cell and extracting the polypeptide of interest
CC
    from the periplasm. The method is useful for preparing a wide variety of
CC
    recombinant polypeptides of interest such as human interferon alpha 2.
CC
    The present sequence is that of a region of the B diminuta gaclss gene
CC
    which was used in the exemplification of the invention.
XX
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     05-MAY-2005 (first entry)
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     24-FEB-2005.
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     12-AUG-2004; 2004WO-EP009067.
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     13-AUG-2003; 2003US-0494914P.
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     Windisch J, Schoergendorfer K, Palma N, Knauseder F, Boehling H;
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XX
DR
     WPI; 2005-182378/19.
DR
     P-PSDB; ADY34498.
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PT
     New expression vector comprising a polynucleotide encoding a fusion
PT
     protein comprising the signal sequence of the gac gene of Pseudomonas
     diminuta and a polypeptide other than gac, useful for producing
PT
PΤ
     polypeptides.
XX
PS
     Example 1; SEQ ID NO 19; 39pp; English.
XX
CC
     The invention provides a process for the efficient and direct production
CC
     of a mature recombinant polypeptide in a prokaryotic host cell. A claimed
CC
     expression vector comprises a polynucleotide encoding a fusion protein
CC
     consisting of the signal sequence ADY34482 of the glutaryl 7-
CC
     aminocephalosporic acid acylase (gac) gene of Pseudomonas diminuta and
CC
     the polypeptide of interest. A prokaryotic host cell transformed with the
CC
     vector is cultured under conditions which cause expression of the
     polynucleotide. Upon formation of the fusion protein, the signal sequence
CC
CC
     is cleaved off and the polypeptide of interest is released into the
     periplasm of the host cell. The expression vector is a plasmid,
CC
     preferably a high copy plasmid. The vector further comprises a
CC
CC
     polynucleotide comprising the promoter region and the ribosomal binding
```

```
site ADY34485 or ADY34486 of the gac gene of P. diminuta. The culturing
CC
     is performed as a multi-stage fermentation process comprising a shake-
CC
     flask step, optionally a pre-culture step, and a main culture step. The
CC
    main culture step is performed in a culture medium comprising a substrate
CC
     for more than 90% of the cultivation time at a substrate concentration
CC
     lower that the saturation constant of the substrate, accompanied by high
CC
     levels of dissolved oxygen concentration, and further accompanied by a
CC
     steadily decreasing specific growth rate of the bacterial host cells, the
CC
     process being performed at a temperature that is lower than the optimum
CC
CC
     temperature for growth of the host cell. The substrate is glycerol or a
CC
     carbohydrate, preferably glucose. The process is favorably used for the
CC
     production of recombinant human interferon-alpha 2B in Escherichia coli.
CC
     The present sequence is that of an expression construct for recombinant
CC
     production of human mature interferon-alpha 2B in E. coli. It comprises
CC
     the P. diminuta signal sequence, promoter and ribosome binding site, and
     the coding sequence ADY34481 for interferon-alpha 2B in which codons have
CC
     been modified to improve expression. The expression construct was
CC
     obtained by a combination of chemical synthesis and PCR amplification.
CC
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  Matches 100; Conservative
                                                                         0;
                               0; Mismatches
                                                0;
                                                    Indels
                                                                  Gaps
           1 ATCCTGGTTCGTACGCGCCCCTACAAGTGGTGATCTAGGGGAACGTTCCGGGGGCGTCG 60
Qу
             110 ATCCTGGTTCGTACGCGCCCTACAAGTGGTGATCTAGGGGAACGTTCCGGGGGCGTCG 169
Db
          61 CTGCAACGCCTCTCCGGATCTGGGTGAGAGGGGAAATCC 100
Qy
             Db
         170 CTGCAACGGCGTCTCCGGATCTGGGTGAGAGGGGAAATCC 209
<!--EndFragment-->
```